

(12) UK Patent Application (19) GB (11) 2 041 962 A

(21) Application No 7942717

(22) Date of filing  
11 Dec 1979

(30) Priority data

(31) 971625

(32) 20 Dec 1978

(33) United States of America  
(US)

(43) Application published  
17 Sep 1980

(51) INT CL<sup>3</sup> C09K 3/34  
C09B 57/00 //  
C07C 50/24

(52) Domestic classification  
C4X 12  
C2C 200 220 221 227  
22Y 313 31Y 338 351  
353 364 36Y 388 43X  
624 635 662 AA UA  
C4P D1G D1R

(56) Documents cited  
None

(58) Field of search  
C4P  
C4X

(71) Applicant  
Timex Corporation  
Waterbury  
Connecticut 06720  
USA

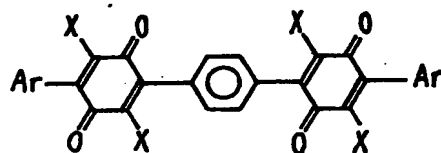
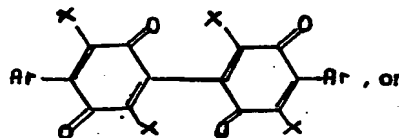
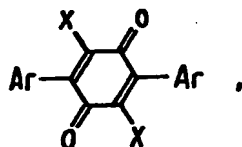
(72) Inventor  
Gene Carl Koch

(74) Agents  
Stevens Hewlett &  
Perkins

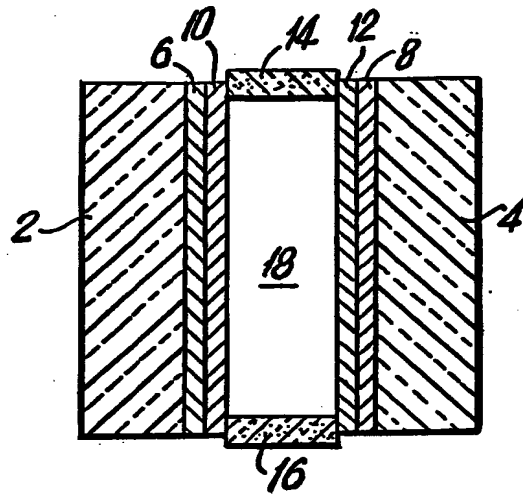
(54) Guest-host liquid crystal composition

(57) Guest coloured material for use in an electro-optic liquid crystal composition has the formula:-

where Ar represents an aromatic (carbocyclic or heterocyclic) ring or ring system in each case, and X is generally a blocking substituent to ensure a linear molecular structure, although in some disclosed materials it is H.



GB 2 041 962 A



## SPECIFICATION

## Electro-optical composition of the guest-host type

## 5 Field of the Invention:

The present invention relates to liquid crystal display devices and, more particularly, to those of the guest-host type in which a dye is admixed with a liquid crystal material.

## Description of the Prior Art:

- 10 A display, particularly one of the liquid crystal type, can be characterized by brightness and contrast performance criteria. It is known to improve these performance criteria by dissolving a guest dichroic dye in a host liquid crystal material. Many dyes have been especially developed for a variety of applications such as dyeing of fabrics, printing of textiles, coloring of plastics, color image formation in photography, etc. To provide the requisite properties, such as hue, solubility, affinity for the substrate, chemical resistance, and compatibility with the medium from 15 which the dye is applied, the molecular structure of the dye is specially designed for each application. Important dye properties required for the present application in liquid crystal displays, include dichroism, high order parameter and solubility.

- Dichroic materials display optical anisotropy, that is to say light whose electric vector 20 component is parallel to a preferred molecular axis of a dichroic molecule will be absorbed more strongly than light whose electric vector components lie along other axes. More specifically, in positive dichroic dyes, such as those treated in the present invention, light which impinges on the dye molecule perpendicular to its long axis is absorbed more strongly than light which impinges on the molecule parallel to its long axis.

- 25 If such a positive dichroic dye is dissolved in a nematic liquid crystalline host, then the long axis of individual dye molecules will be constrained to align with the nematic material. When the nematic host material is aligned by either an external electric field or preselected boundary conditions in the display, the dye molecules are constrained to adopt the alignment of the liquid crystal. This effect is put to use in guest-host displays by using boundary conditions, such as by 30 molecular ordering induced by an angularly deposited silicon oxide coating, to initially align the dye long axes in one direction and then imposing an electric field so as to rotate the dye long axes in a perpendicular direction, thereby altering the light absorbing properties of the display.

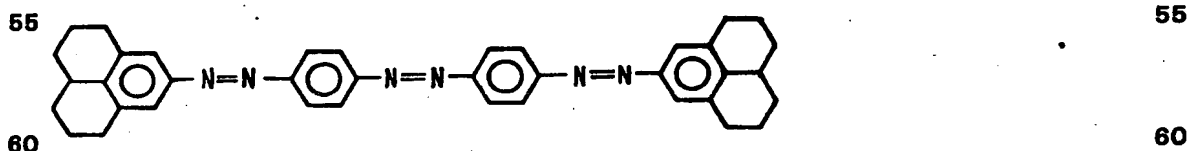
In order for this guest-host device to perform properly, the guest dye molecules must align well with the host nematic liquid crystal material.

- 35 The efficiency of this alignment is commonly determined aligning a guest-host (dye-nematic liquid crystal) solution with an electric field so that the absorption of polarized light beams whose electric vectors lie parallel and perpendicular to the long dye axis, respectively, can be measured and compared. The alignment efficiency can be expressed as an order parameter,  $S$ , which is given by:

$$40 \quad S = \frac{A_{11} - A_1}{A_{11} + A_1}$$

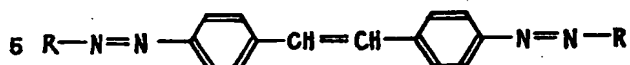
- 45 where  $A_{11}$  and  $A_1$  are, respectively, the absorbance for light polarized with electric vectors along the long axis and the absorbance for light polarized with electric vectors perpendicular to the long axis. The order parameter, however, is not solely dependent on the efficiency of the alignment of the guest dye with the nematic material but also depends on the intrinsic optical anisotropy of the dye molecules and the order parameter of the host nematic material itself.

- 50 Experience with dichroic dyes for use in guest-host display devices has shown that generally the longer and more rodlike dye molecules are more likely to have higher order parameters. For example, one type of dye capable of sufficiently high order parameters to yield displays with good viewing properties are azo dyes, for instance:



As described in the Aftergut et al patent, U.S. 4,116,861, issued September 26, 1978, another dye which might exhibit a sufficiently high order parameter is a stilbene-derived dichroic dye, for example, of the following structure:

65



5

10 Solubility of the dichroic dye in the liquid crystal must be sufficiently high so that thin layers, for example, of ten to fifteen microns, have adequate light absorption in one of the oriented states.

10

15 Liquid crystal-dye compositions utilized in electrooptical displays are discussed in the Goldmacker et al patent, U.S. 3,499,702 issued March 10, 1970; Castellano patent, U.S. 3,579,044 issued August 3, 1971; Castellano patent, U.S. 3,703,329 issued November 21, 1972; Hatfield et al patent, U.S. 3,837,730 issued September 24, 1974; Moriyama et al patent, U.S. 3,864,022 issued February 4, 1975; Moriyama et al patent U.S. 3,960,750 issued September 1, 1976; Moriyama et al patent, U.S. 3,960,751 issued September 1, 1976; Constant et al patent U.S. 4,032,219 issued June 28, 1977 and Bloom patent, U.S. 4,032,470 issued June 28, 1977.

15

20

## 20 SUMMARY OF THE INVENTION

It is an object of the invention to provide an electro-optical composition comprising a host liquid crystal and a guest dichroic dye admixed therein, the dye molecules being long and rodlike so as to exhibit a high order parameter and being adequately soluble in commonly used liquid crystals or mixtures thereof for display purposes.

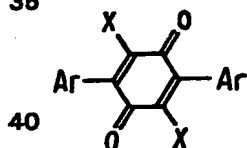
25

It is another object of the invention to provide an improved electro-optical display by virtue of the incorporation of the aforementioned electro-optical composition therein.

In the electro-optical composition of the invention, the host liquid crystals may include conventional and well known materials such as nematic and cholesteric liquid crystals and their mixtures while the guest dichroic dye is characterized generally as a linear noncondensed aromatic compound which includes a quinonoid ring as the chromophore (that is, as the color-generating group). The inventive dichroic dye is selected from a compound represented by the formula:

30

35



35

40

45 where Ar is a mono or polycyclic aromatic ring system which may include one or more additional quinonoid rings in the paraposition and/or hetero atoms and where X is a blocking substituent such as Cl, Br, F, H, an acryloxy group or alkoxy group to ensure linear characteristics.

45

## BRIEF DESCRIPTIONS OF THE DRAWINGS

50 The Figure is a cross-sectional view of an electro-optic device embodying the invention.

50

## DESCRIPTION OF THE PREFERRED EMBODIMENTS

A typical electro-optical display device embodying the invention is illustrated schematically in the Figure and includes two glass or other transparent substrates 2 and 4 having conductive oxide electrodes 6 and 8 thereon. For example, electrodes 6 and 8 may comprise tin oxide, indium oxide or admixtures thereof. Evaporated or otherwise applied onto the oxide electrodes are alignment-promoting layer 10 and 12, for example, thin silicon oxide layers evaporated at an angle of 60°. Glass frit spacers 14 and 16 maintain the substrates 2 and 4 apart to complete the cell. The cell is baked at an elevated temperature to melt the glass frit and seal the cell except for a fill port. The electro-optical composition of the invention is then introduced into the space 18 between the substrates and the fill port sealed. Of course, electrodes 6 and 8 are connected to suitable circuitry to generate the desired electric field across the electrodes to rotate the host liquid crystal molecules and thus the guest dichroic dye molecules to vary the light absorption characteristics of the cell.

55

60

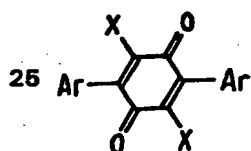
65 The electro-optical composition of the present invention comprises a host liquid crystal and

65

guest dichroic dye admixed together. Preferably the liquid crystal is a nematic material exhibiting strong positive dielectric anisotropy but other liquid crystals, particularly of the cholesteric type, may be employed depending upon the electro-optic effect to be utilized. Low melting point nematic liquid crystals, such as p-alkoxybenzylidene-p'-alkylanilines and their mixtures with other liquid crystals, and mixtures of p-alkoxy- or acryloxybenzylidene-p'-cyanoanilines, having a transition temperature range that includes room temperature, are suitable. Other nematic liquid crystals useful in the invention include, but are not limited to, esters available from Merck Company and Hoffmann LaRoche Company, biphenyls from BDH Ltd., phenyl cyclohexanes and azoxy compounds available from Merck Company as well as others. Suitable cholesteric liquid crystals can be made by incorporating optically active compounds in any of the above nematic liquid crystals.

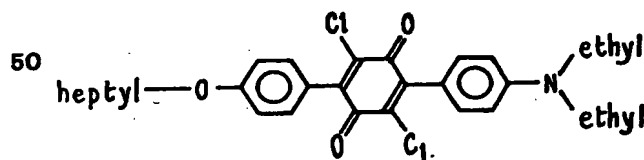
In preparing an electro-optic display device of the invention, the liquid crystal materials should be rigorously purified to remove ionic and nonionic impurities which may react to degrade the liquid crystal compounds by decomposition, transubstitution reactions and the like. For commercially acceptable devices, the liquid crystal compounds should be purified so that their resistivity is  $10^{11}$  ohm-cm or higher.

An important feature of the present invention is that the guest dichroic dye is characterized as a linear, noncondensed aromatic compound including at least one quinonoid ring in the molecular structure as the color generating group (or chromophore), the compound being represented by the formula:



where Ar represents a mono or polycyclic aromatic ring system which may include one or more additional quinonoid rings in the para position and which may also include hetero atoms and where X is a blocking substituent to ensure a linear molecular structure. X may include Cl, Br, F, H, and acyloxy group or alkoxy group for this purpose. Generally, the dye compound represented by the above formula may be referred to as 2,5-diaryl-1,4 benzoquinone. The advantageous properties of dyes of this type are attributable to their long, rod-like structure which results in a high order parameter and thus in improved contrast and to the quinonoid ring as the color-generating group. These dyes are generally soluble in all the commonly used nematic liquid crystals listed in part hereinabove, and impart a variety of colors, including yellow, red, green, purple, etc. to a solution thereof (depending on the specific dye compound utilized) when incident light impinges on the dye molecules perpendicular to their long axes. On the other hand, the solution appears colorless when incident light impinges parallel to their long axes.

An exemplary dichroic dye of the invention where Ar does not include additional quinonoid rings is 2-(4-heptyloxyphenyl)-5-{4-(N,N-diethylamino) phenyl}-3,6-dichlorobenzo-4-quinone represented by the formula:



An exemplary synthesis of this dichroic dye involves the following two-step process including:

**Step A: 2-(4-Heptyloxyphenyl)-3,6-dichlorobenzo-4-quinone:**

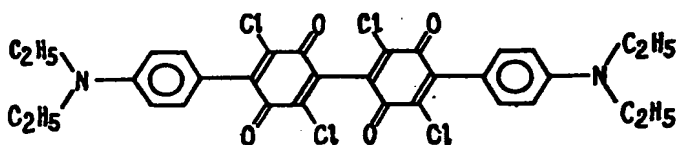
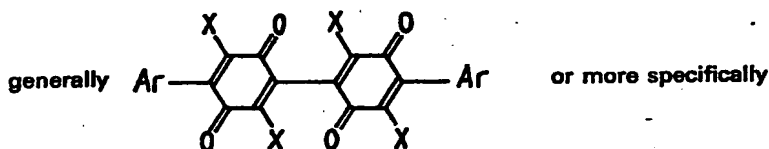
34g (.16 moles) of 4-heptyloxyaniline is dissolved in 51 ml of concentrated hydrochloric acid. The solution is diluted with 25 ml of water and chilled to 5°C in an ice bath. The solution is treated with a paste consisting of 12.9g sodium nitrite in 36 ml water with vigorous stirring. When diazotization is complete it is added simultaneously with 28.4g of sodium acetate in 50 ml of water to a stirred solution of 25g (.14 moles) of 2,5 dichlorobenzo-4-quinone in 780 ml methanol and 210 ml diethyl ether with an additional 25 ml water added to rinse. This mixture

is stirred for 2 hours, then 400 ml of solvent is stripped off and 200 ml boiling water added. The solution is brought to a boil and then filtered hot. On cooling the filtrate precipitated orange crystals are collected by filtration. The precipitated product gives only one spot by TLC. The red tap which is collected in the first filtration proved to be largely product material mixed with some of the diheptyloxyphenyl quinone. This material is purified by recrystallization from ethanol/water.

**Step B: 2-(4-Heptyloxyphenyl)-5-(4-N,N-diethylamino)phenyl-3,6-dichlorobenzo 4-quinone:**

16g (.089 moles) of N,N-diethyl-p-phenylenediamine is added to 20 ml concentrated hydrochloric acid and chilled in an ice/salt bath to  $-5^{\circ}\text{C}$ , then diazotized with a paste consisting of 7g sodium nitrite in 20 ml water. When the diazotization is complete sufficient sodium acetate is added to buffer the solution to neutrality. The buffered diazo solution is added to a vigorously stirred solution of 4g (.011 moles) of the Heptyloxyphenyl quinone (Compound A) in 400 ml ethanol and 200 ml of diethyl ether. The red solution immediately begins to darken and bubbles of nitrogen are formed. The reaction mixture is allowed to stand overnight and then is diluted with an equal volume of water. The solution is stripped of alcohol and ether on a vacuum and then extracted with methylene chloride. The extract is dried and stripped to give the crude product. A gross purification is carried out on an open silica gel column using 5% diethyl ether in hexane as the elutant. The product material obtained from this step is further purified using a Waters Associates Pres 500 preparative liquid chromatograph with the standard Pred Pak silica cartridge and a 2% ethyl ether /98% hexane elutant. Compound purity is monitored by thin layer chromatography. Melting point  $-114^{\circ}\text{C}$ .

An exemplary dichroic dye of the invention where Ar includes an additional quinonoid ring in the para position is 5,5'-di 4-(N,N-diethylamino)phenyl-3,3', 6,6'-tetrachloro-2,2'-bibenzo-4-quinonyl having the formula:



A typical synthesis of this compound also requires two steps as follows:

**Step A: 3,3', 6,6'-tetrachloro-2,2'-bibenzo-4-quinonyl:**

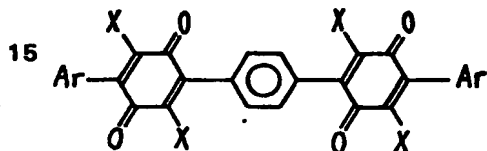
10g. of 2,5-dichlorohydroquinone dimethyl ether is dissolved in 125 ml. of glacial acetic acid with stirring. 15 ml. of 6 N sodium dichromate solution is added dropwise over 10 minutes. The dichromate color is discharged immediately and a green color develops. The solution becomes very warm. The mixture is stirred for ten more minutes, then poured onto 100g. of an ice/water mixture. The red precipitate formed is collected by filtration and washed with water. The product diquinone is recrystallized from a minimum of ethyl alcohol to yield long red needles. (Melting point  $-129^{\circ}\text{C}$ , with considerable sublimation and pretransitional polymorphic behavior.)

**Step B: 5,5'-Di 4-(N,N-diethylamino)phenyl-3,3', 6,6'-tetrachloro-2,2'-bibenzo-4-quinonyl:**

3.1 g. (.028 moles) of N,N-diethylphenylenediamine is dropped into 5mL. of hydrochloric acid. The amine is diazotized by addition of a paste of 1.33 g. sodium nitrite in 5mL. water with vigorous stirring. When the diazotisation is complete, sufficient sodium acetate is added to buffer the solution to a pH of approximately 7. The buffered diazo solution is added immediately to a solution of 4.9 g. (.014) moles of the diquinone (compound A) in 300 mL. ethanol and 250 mL. diethyl ether. The reaction mixture begins to darken immediately and after 30 minutes the solvent is stripped from the mixture under vacuum. Prolonged exposure of the product to

the reaction conditions leads to deterioration and reduced yield. The residue remaining after the solvent is removed is dissolved in 50 mL. methylene chloride and passed through silica gel column with methylene chloride elutant so as to remove tars and other gross contaminants. The crude blue product obtained from this step is further purified using a Waters Assoc. Prep 500 preparative liquid chromatography with the standard Prep Pak silica cartridge and a 15% diethyl ether/85% hexane elutant. Compound purity is monitored by thin layer chromatography. The yield is .3g. of highly purified dye (3.9% yield). Melting point  $-128^{\circ} - 130^{\circ}\text{C}$  with prior softening.

Another exemplary dichroic dye of the invention where Ar includes an additional quinonoid ring in the paraposition may be referred to as 1,4 di(5 aryl benzo-4-quinon-2yl) benzene having the formula:



where Ar is a mono or polycyclic aromatic ring system with or without hereto atoms and X is a blocking substituent as described hereinabove.

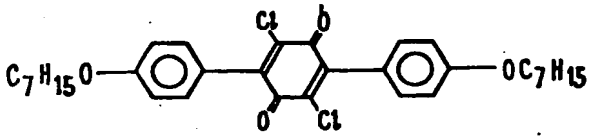
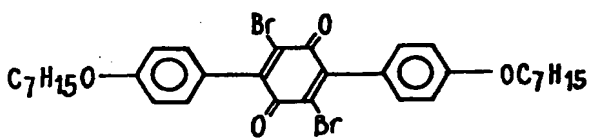
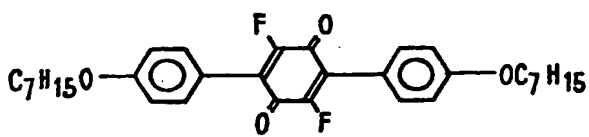
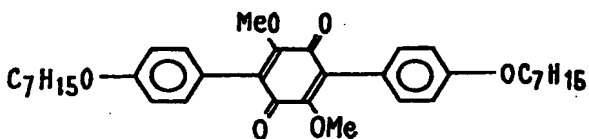
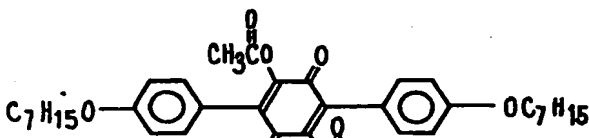
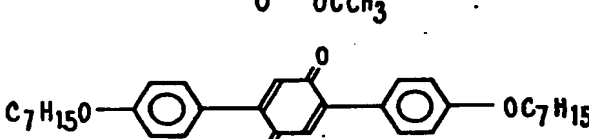
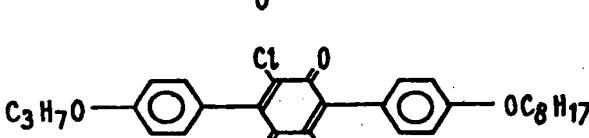
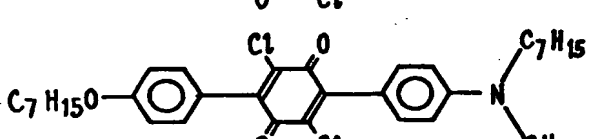
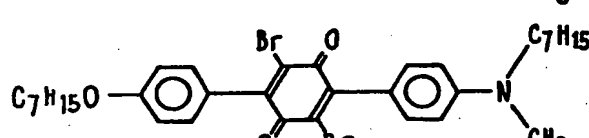
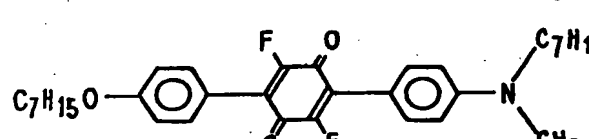
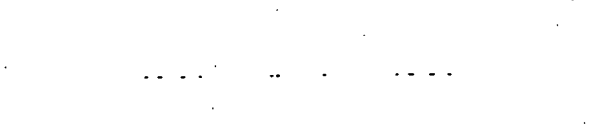

The dichroic dyes of the invention can be admixed with nematic or other liquid crystals to impart a characteristic color thereto under certain alignment conditions. The exact amount of dye admixed in the liquid crystal depends on the solubility of the dye in the liquid crystal and also on the strength of the color desired. In general, an electro-optical composition of the invention will include from about 1% to about 4% by weight of the dichroic dye. For example: a working display was prepared by dissolution of 3% by weight of 2,5-di(4-heptyloxy phenyl)-3,6 dichlorobenzo-4-quinone in a liquid crystalline mixture composed of 42% 4-heptyl-4'-cyanobiphenyl, 23% 4-octyloxy-4'-cyano biphenyl, 19% 4-pentyloxy-4'-cyanobiphenyl and 16% 4-heptyloxy-4'-cyanobiphenyl.

A display cell such as previously described with a spacing between substrates of  $12\mu$  was filled with material and operated at 3 volts (rms). This gave a working device with white digits on a yellow background.

Similarly a display mixture containing 3% 2-(4-heptyloxyphenyl) -5-(4-(N,N-diethyl amino) phenyl)-3,6-dichlorobenzo-4-quinone in the above liquid crystal composition was filled into a similar display cell and operated at 3 volts (rms) to yield a display with white digits on a green background.

The following table is provided to illustrate in greater detail dichroic dyes of the invention. These examples are for purposes of illustration rather than limitation.

TABLE  
1

5		5
10		10
15		15
20		20
25		25
30		30
35		35
40		40
45		45
50		50
55		55
60		60
65		65



**TABLE (Cont.)**

11.		5
12.		10
13.		20
14.		25
15.		30
16.		40
17.		45
18.		50
19.		55
		60

TABLE (Cont.)

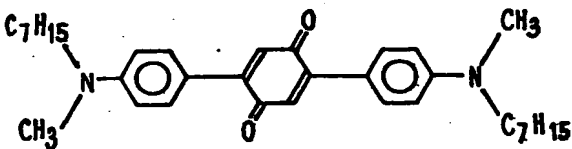
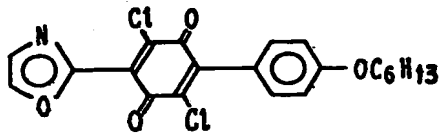
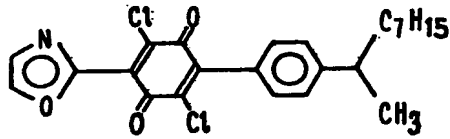
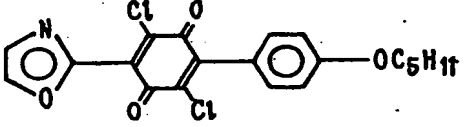
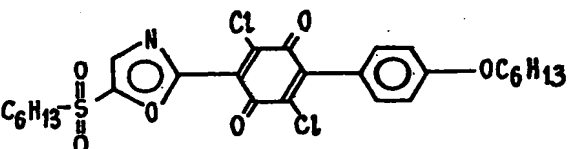
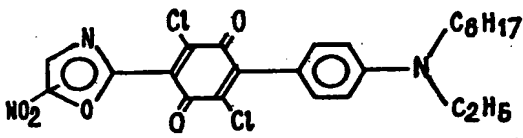
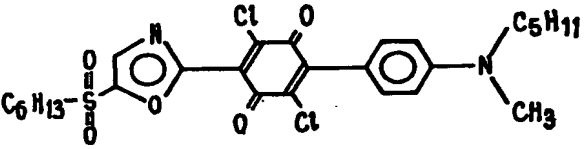
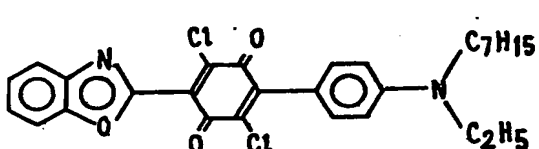
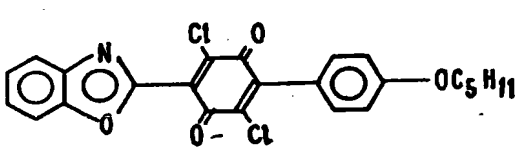
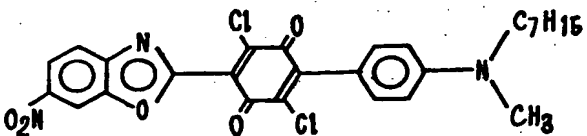
20.		5
21.		10
22.		15
23.		20
24.		25
25.		30
26.		35
27.		40
28.		45
29.		50

TABLE (Cont.)

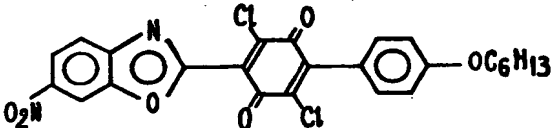
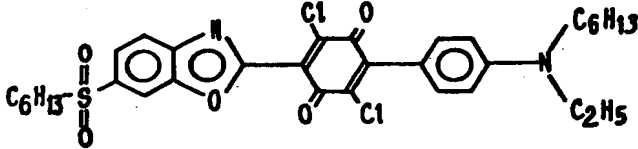
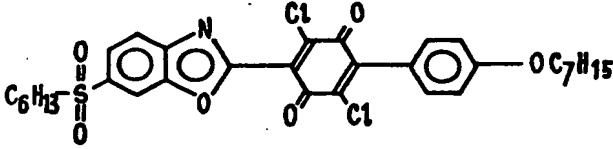
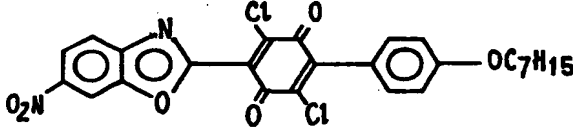
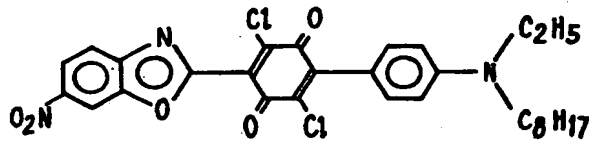
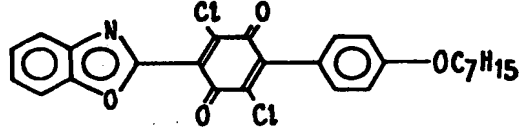
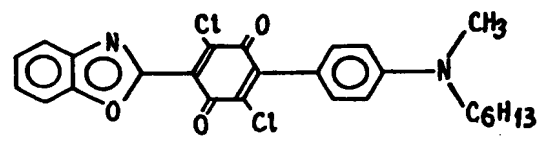
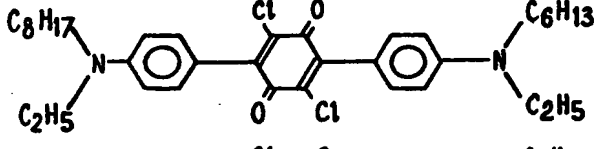
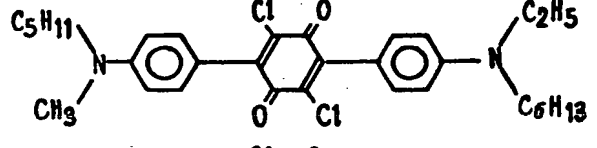
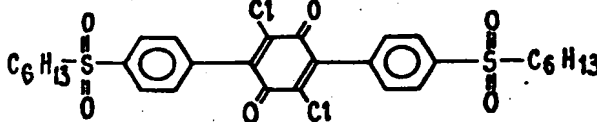
30.		5
31.		10
32.		15
33.		20
34.		25
35.		30
36.		35
37.		40
38.		45
39.		50

TABLE (Cont.)

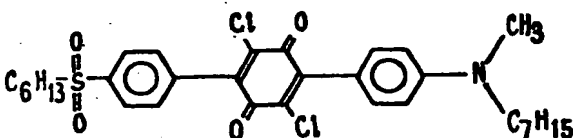
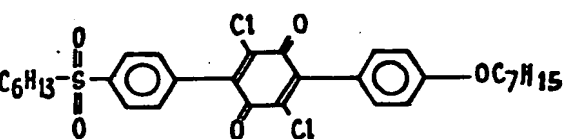
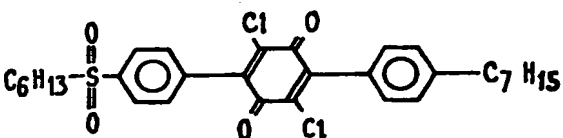
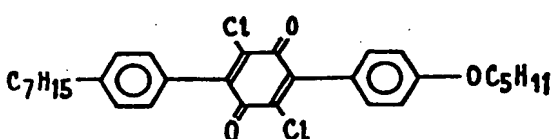
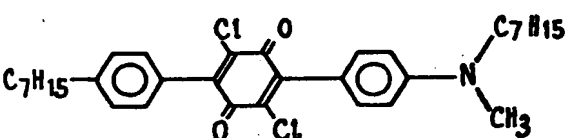
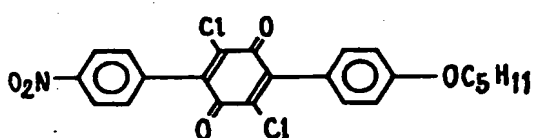
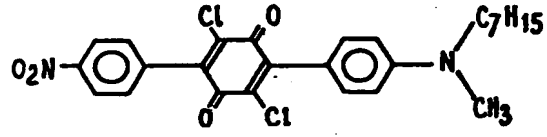
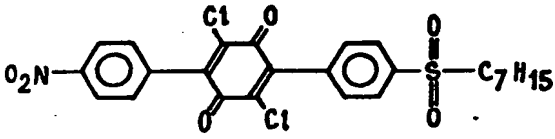
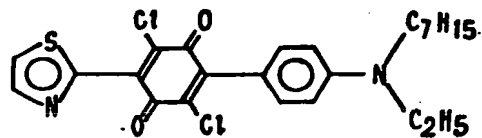
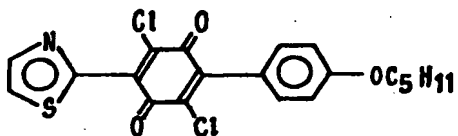
40.		5
41.		10
42.		15
43.		20
44.		25
45.		30
46.		35
47.		40
48.		45
49.		50

TABLE (Cont.)

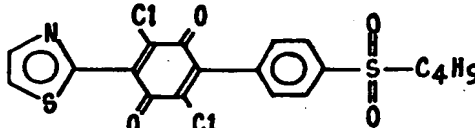
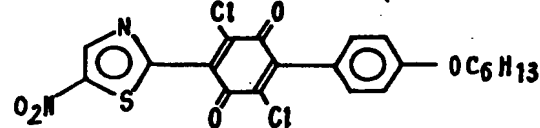
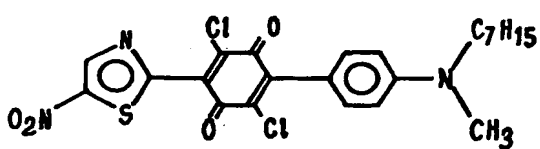
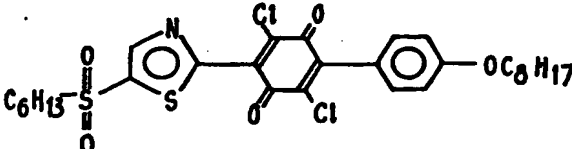
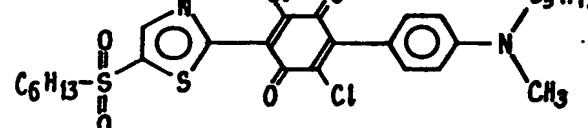
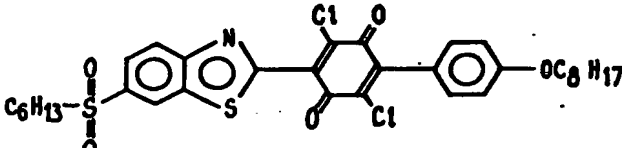
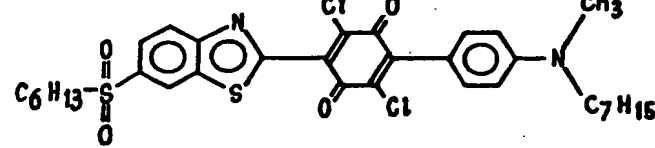
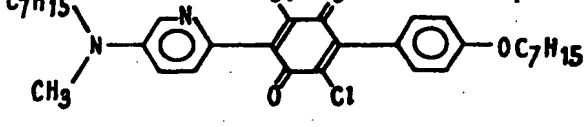
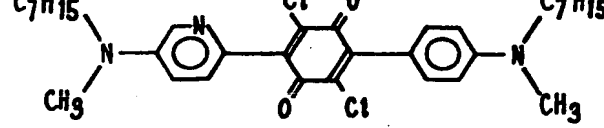
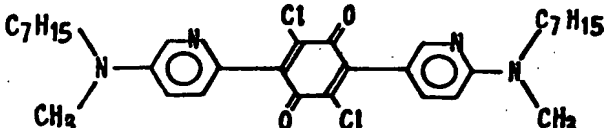
50.		5
51.		10
52.		15
53.		20
54.		25
55.		30
56.		35
57.		40
58.		45
59.		50

TABLE (Cont.)

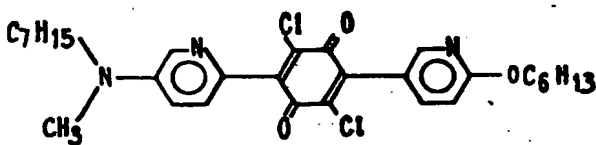
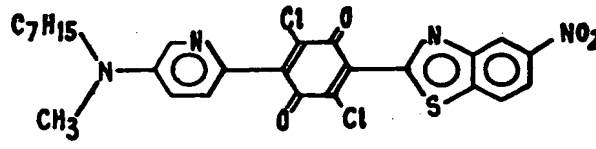
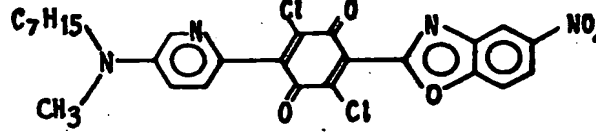
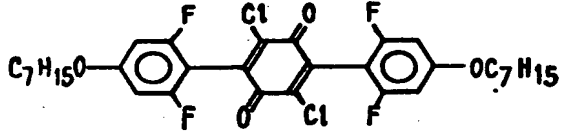
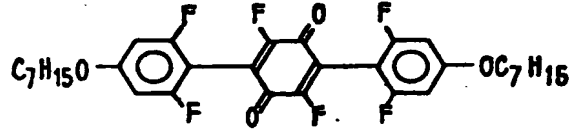
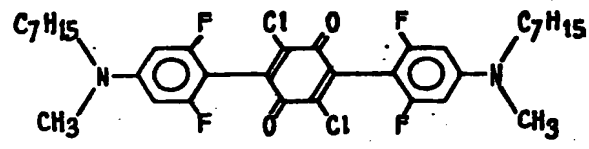
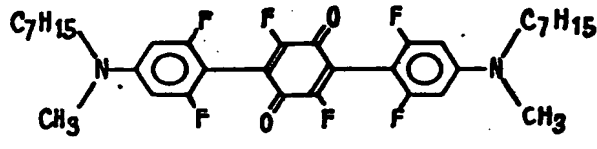
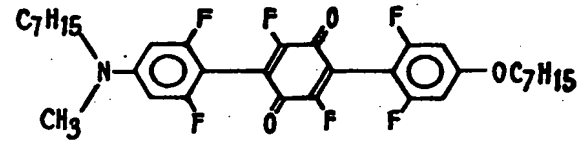
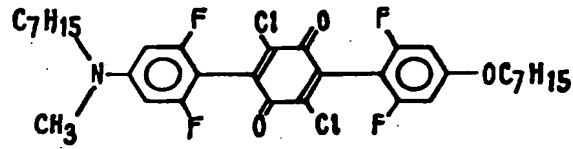
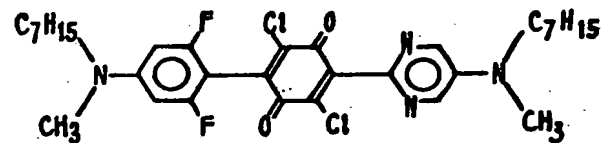
60.		5
61.		10
62.		15
63.		25
64.		30
65.		35
66.		40
67.		45
68.		50
69.		55

TABLE (Cont.)

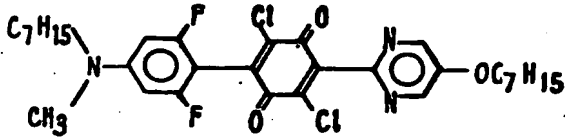
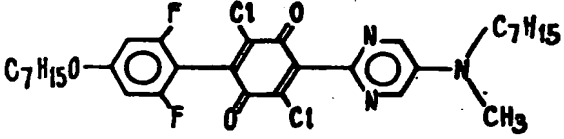
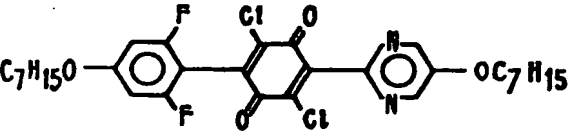
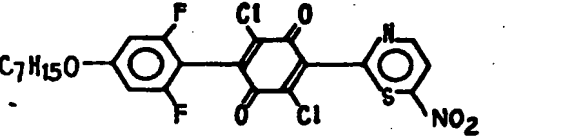
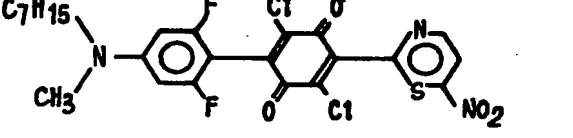
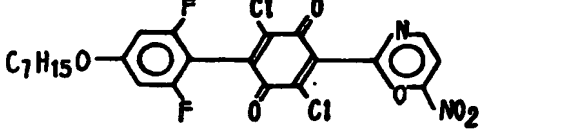
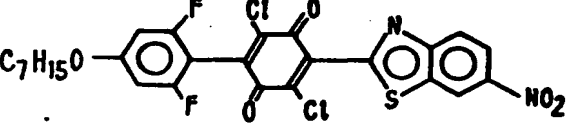
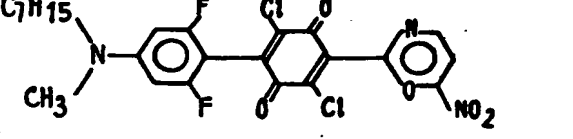
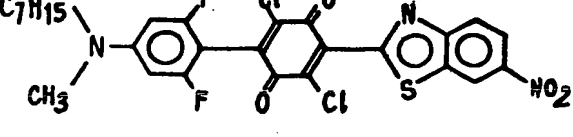
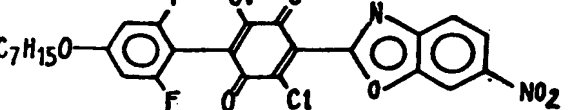
5	70.		5
10	71.		10
15	72.		15
20	73.		20
25	74.		25
30	75.		30
35	76.		35
40	77.		40
45	78.		45
50	79.		50

TABLE (Cont.)

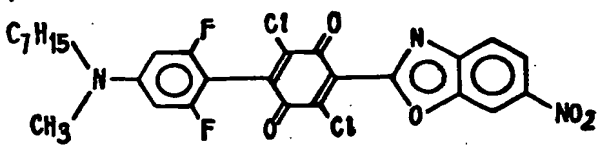
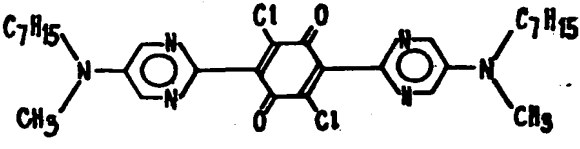
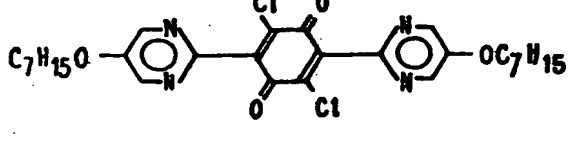
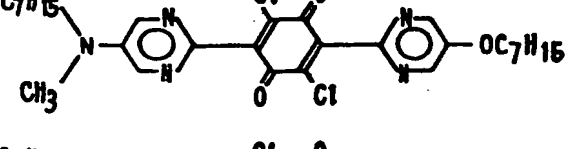
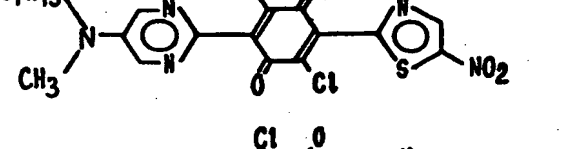
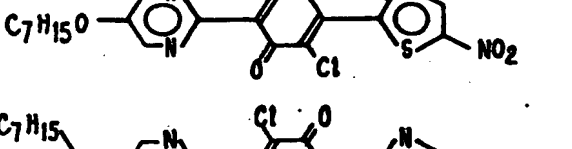
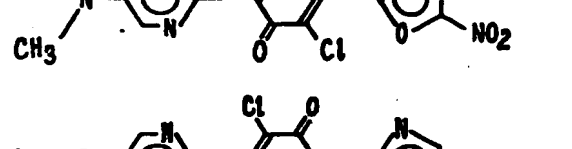
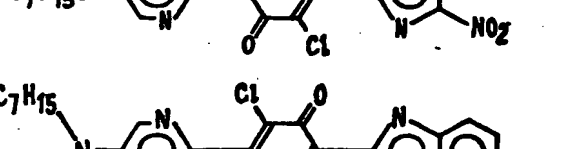
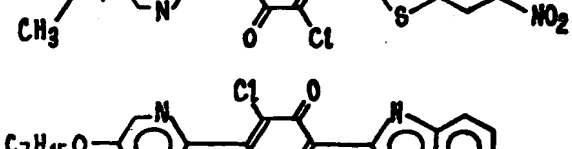
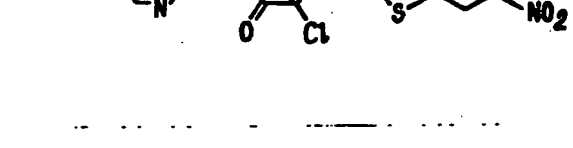
80.		5
81.		10
82.		15
83.		20
84.		25
85.		30
86.		35
87.		40
88.		45
89.		50



TABLE (Cont.)

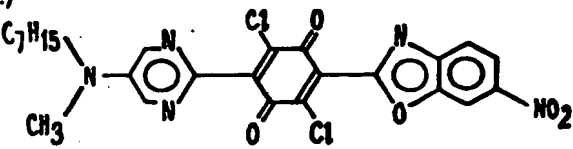
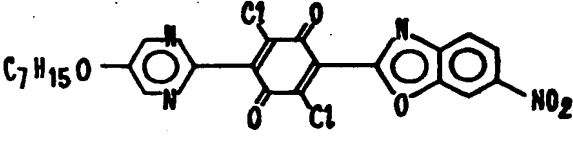
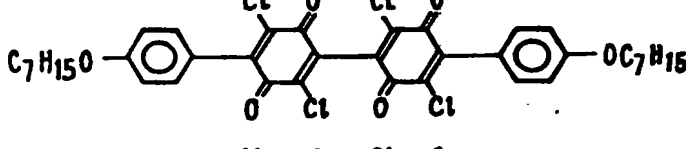
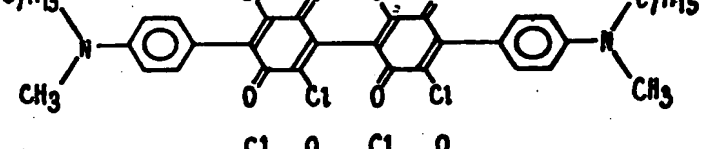
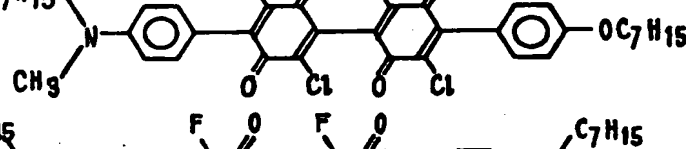
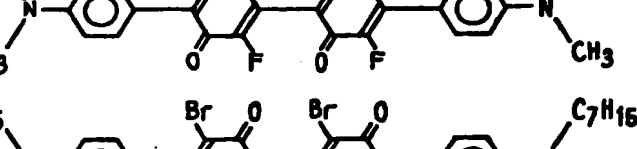
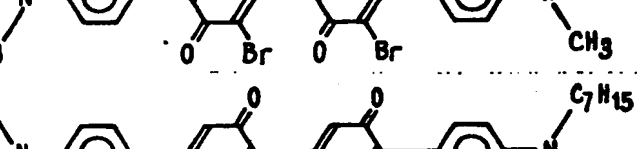
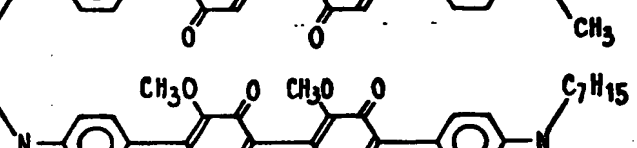
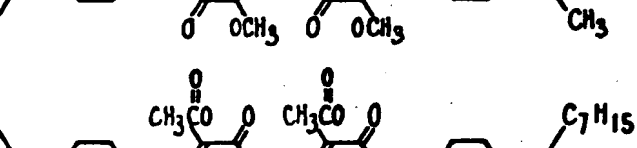
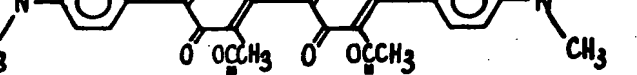
90.		5
91.		10
92.		15
93.		20
94.		25
95.		30
96.		35
97.		40
98.		45
99.		50

TABLE (Cont.)

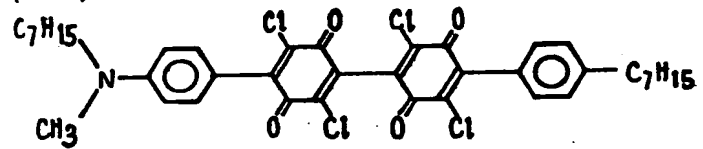
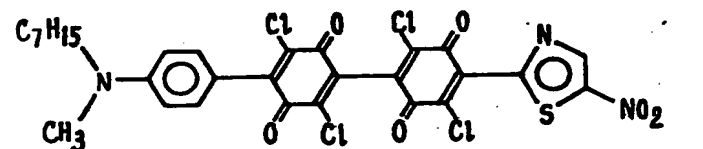
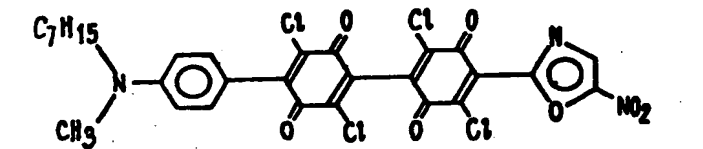
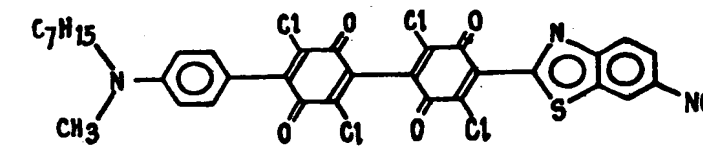
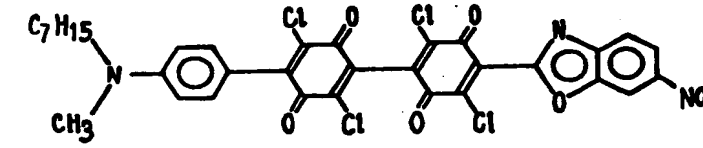
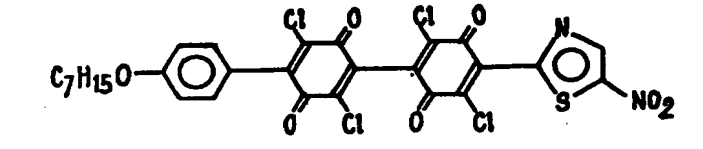
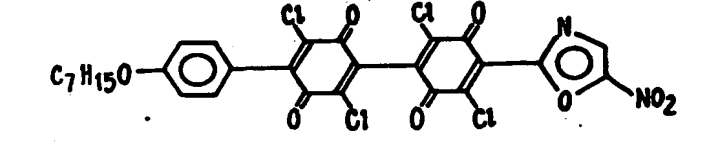
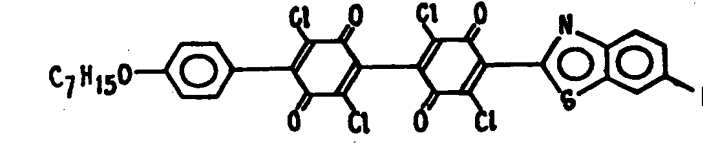
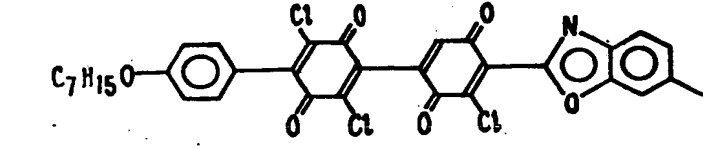
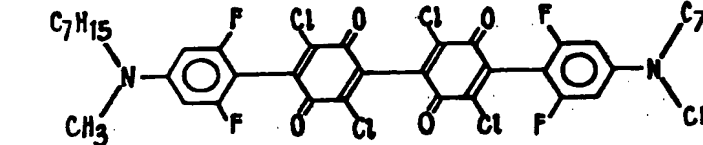
100.		5
101.		10
102.		15
103.		20
104.		25
105.		30
106.		35
107.		40
108.		45
109.		50

TABLE (Cont.)

5	110.		5
10	111.		10
15	112.		15
20	113.		20
25	114.		25
30	115.		30
35	116.		35
40	117.		40
45	118.		45
50	119.		50
55			55
60			60

TABLE (Cont.)

120.		5
121.		10
122.		15
123.		20
124.		25
125.		30
126.		35
127.		40
128.		45
129.		50

TABLE (Cont.)

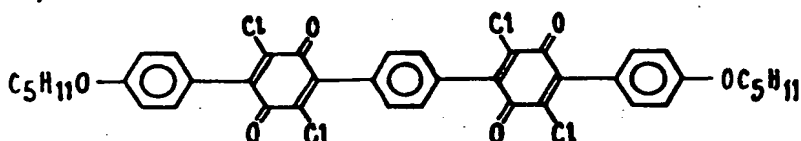
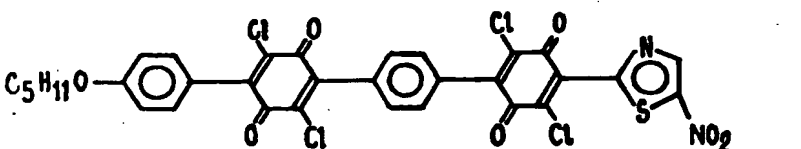
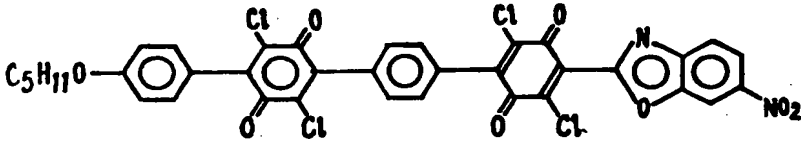
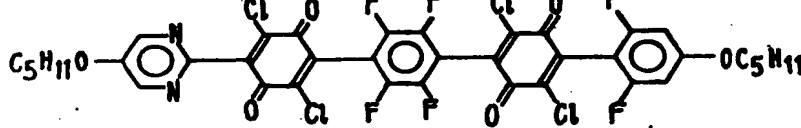
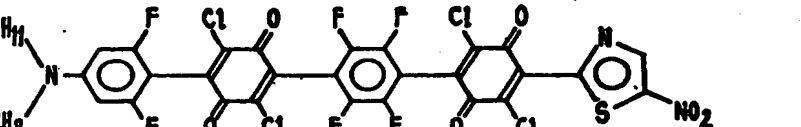
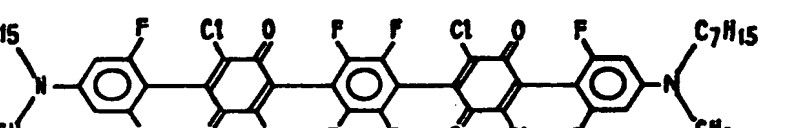
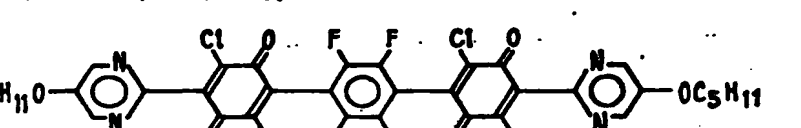
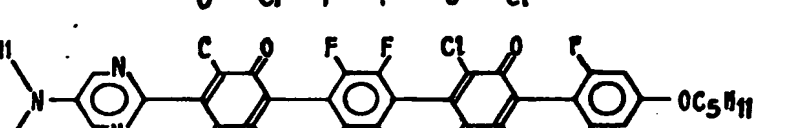
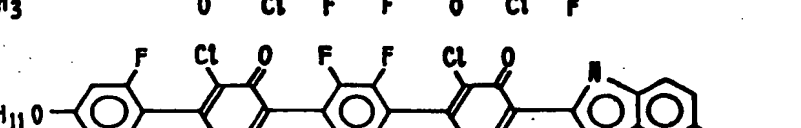
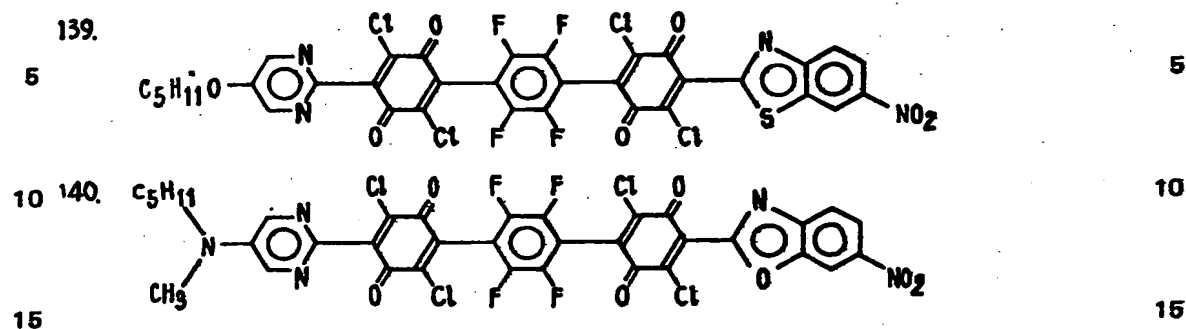
5	130.		5
10	131.		10
15	132.		15
20	133.		20
25	134.		25
30	135.		30
35	136.		35
40	137.		40
45	138.		45
50			50
55			55

TABLE (cont.)



While there has been described what is considered to be the preferred embodiments of the invention, other modifications will occur to those skilled in the art and it is desired to cover in the appended claims all such modifications as fall within the true spirit and scope of the invention.

## CLAIMS

**CLAIMS**

1. An electro-optical composition comprising a host liquid crystal admixed with a linear, noncondensed aromatic guest dichroic dye of the chemical formula:

25

25



**35** where Ar represents an aromatic ring system which may include additional quinonoid rings and **35**  
**X** is a blocking substituent to ensure a linear molecular structure.

2. An electro-optical composition comprising a host liquid crystal admixed with a linear, noncondensed aromatic guest dichroic dye of the chemical formula:



where Ar represents an aromatic ring system and X is a blocking substituent to ensure a linear molecular structure.

3. An electro-optical composition comprising a host liquid crystal admixed with a linear, noncondensed aromatic guest dichroic dye of the chemical formula:



where Ar represents an aromatic ring system and X is a blocking substituent to ensure a linear molecular structure.

65 4. The composition of any one of claims 1 to 3 wherein each aromatic ring system, Ar, of 65

the dichroic dye may include hetero atoms.

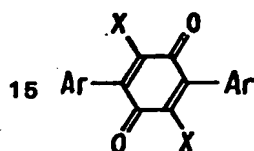
5. The composition of any one of claims 1 to 4 wherein the substituent X, of the dichroic dye can be H, Cl, Br, F, an Acyloxy group or an alkoxy group.

6. The composition of any one of claims 1 to 5, wherein the dichroic dye is present in an amount of from 1% to 4% by weight.

7. In an electro-optical display device comprising two spaced transparent substrates whose facing surfaces have been coated with transparent electrodes, the improvement comprising an electro-optical composition disposed between the substrates comprising a host liquid crystal admixed with a linear, noncondensed aromatic guest dichroic dye of the chemical formula:

10

10



15

20 where Ar represents an aromatic ring system which may include additional quinonoid rings and X is a blocking substituent to ensure a linear molecular structure.

20

8. The display device of claim 7, wherein the electro-optical composition is as defined in any one of claims 1 to 6.

Printed for Her Majesty's Stationery Office by Burgess & Son (Abingdon) Ltd.—1980.

Published at The Patent Office, 25 Southampton Buildings, London, WC2A 1AY, from which copies may be obtained.

**This Page is Inserted by IFW Indexing and Scanning  
Operations and is not part of the Official Record**

**BEST AVAILABLE IMAGES**

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☐ BLACK BORDERS
- ☐ IMAGE CUT OFF AT TOP, BOTTOM OR SIDES
- ☐ FADED TEXT OR DRAWING
- ☒ BLURRED OR ILLEGIBLE TEXT OR DRAWING
- ☐ SKEWED/SLANTED IMAGES
- ☐ COLOR OR BLACK AND WHITE PHOTOGRAPHS
- ☐ GRAY SCALE DOCUMENTS
- ☒ LINES OR MARKS ON ORIGINAL DOCUMENT
- ☐ REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY
- ☐ OTHER: \_\_\_\_\_

**IMAGES ARE BEST AVAILABLE COPY.**

**As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.**